PCT

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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's	or age	ent's file reference	T		
R1562-PCT			FOR FURTHER AC		ication of Transmittal of International ry Examination Report (Form PCT/IPEA/416)
		ication No.	International filing date (d	dav/month/vear)	Priority date (day/month/year)
PCT/BE			19/06/2000	. 	17/06/1999
		ent Classification (IPC) or na	tional classification and IPC	>	
C07K1/0					
Applicant					
UNIVER	SITE	IT GENT et al.			
			instice report has been	propared by this In	tornational Proliminary Examining Authority
1. This	interna s trans	ational preliminary exam smitted to the applicant a	according to Article 36.	prepared by this in	ternational Preliminary Examining Authority
2. This	REPC	PRT consists of a total of	6 sheets, including this	cover sheet.	
	This re been a	eport is also accompanie Imended and are the bas	d by ANNEXES, i.e. she sis for this report and/or	ets of the descript sheets containing	ion, claims and/or drawings which have rectifications made before this Authority
	see R	ule 70.16 and Section 6	07 of the Administrative	Instructions under	the PCT).
Thes	e ann	exes consist of a total of	sheets.		
3. This	report	contains indications rela	ating to the following iten	ns:	
1	\boxtimes	Basis of the report			
11		Priority			
18	\boxtimes	Non-establishment of o	pinion with regard to no	velty, inventive ste	p and industrial applicability
IV		•			
\ \ \	\boxtimes	Reasoned statement u	nder Article 35(2) with re ons suporting such state	egard to novelty, in ement	ventive step or industrial applicability;
VI		Certain documents cite			
VII		Certain defects in the in	nternational application		
VIII	\boxtimes		n the international applic	cation	
Date of su	bmissio	on of the demand		Date of completion	of this report
10/01/2001				25.10.2001	
Name and	l mailin	g address of the internationa	al	Authorized officer	- croft a
	y exam	ining authority:			is and a second control of the second contro
9)		opean Patent Office - P.B. 5 2280 HV Rijswijk - Pays Ba:		Masturzo, P	(1848) (1848) (1848) (1848) (1848)
اربع	Tel.	+31 70 340 - 2040 Tx: 31 6			1 Tan 19 3 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Fax: +31 70 340 - 3016				Telephone No. +31	70 340 2275

INTERNATIONAL PRELIMINARY **EXAMINATION REPORT**

International application No. PCT/BE00/00066

I. Basis of the report

•	ai	. With regard to the elements of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)): Description, pages:					
	1-	52	as originally filed				
	C	laims, No.:					
	1-	35	as originally filed				
	Dr	awings, sheets:					
	1/2	2-2/2	as originally filed				
2	. Wi lan	th regard to the lang guage in which the i	uage, all the elements marked above were available or furnished to this Authority in the nternational application was filed, unless otherwise indicated under this item.				
	Th	ese elements were a	vailable or furnished to this Authority in the following language: , which is:				
		the language of a t	ranslation furnished for the purposes of the international search (under Rule 23.1(b)).				
		the language of pu	blication of the international application (under Rule 48.3(b)).				
		the language of a t 55.2 and/or 55.3).	ranslation furnished for the purposes of international preliminary examination (under Rule				
3.	Wit	h regard to any nucl ernational preliminary	eotide and/or amino acid sequence disclosed in the international application, the examination was carried out on the basis of the sequence listing:				
		contained in the inte	ernational application in written form.				
			ne international application in computer readable form.				
			ently to this Authority in written form.				
			ntly to this Authority in computer readable form.				
The statement that the subsequently furnished written sequence listing does not go beyond the disting the international application as filed has been furnished.							
		The statement that listing has been furn	the information recorded in computer readable form is identical to the written sequence nished.				
4.	The	amendments have r	esulted in the cancellation of:				
		the description,	pages:				
		the claims,	Nos.:				

4.



International application No. PCT/BE00/00066

		the drawings,	sheets:					
5.		This report has been considered to go be	established	as if (some o	of) the amendr d (Rule 70.2(d	nents had not b ;)):	een made, sind	ce they have been
		(Any replacement st report.)	neet containii	ng such amer	ndments must	be referred to u	ınder item 1 an	d annexed to this
6.	Add	litional observations,	if necessary:	;				
111.	Not	n-establishment of c	pinion with	regard to no	velty, inventi	ve step and inc	dustrial applic	ability
	The	e questions whether the rious), or to be industi	ne claimed in	vention appe	ars to be nove	el, to involve an		
		the entire internation	nal applicatio	n.				
	×	claims Nos. 1-35.						
be	cau	se:						
		the said international not require an interr	al application national prelin	, or the said o minary examin	claims Nos. re nation (<i>specif</i> y	late to the follow /):	wing subject ma	atter which does
	×	the description, clai unclear that no mea see separate shee	ningful opini	gs (<i>indicate p</i> on could be fo	particular elem ormed (specify	ents below) or s /):	said claims Nos	s. 1-35 are so
		the claims, or said could be formed.	elaims Nos.	are so inadeq	uately suppor	ted by the desci	ription that no r	neaningful opinior
		no international sea	rch report ha	as been estab	lished for the	said claims Nos	3. .	
2.	and	neaningful internatior d/or amino acid seque tructions:	al preliminar ence listing to	y examination o comply with	n cannot be ca the standard	arried out due to provided for in A	the failure of t Annex C of the	he nucleotide Administrative
		the written form has	s not been fu	rnished or do	es not comply	with the standa	ard.	
		the computer reada						ı .
٧	. Re	easoned statement u ations and explanat	inder Article ions suppoi	35(2) with re	egard to nove atement	elty, inventive s	step or indust	rial applicability;
1	. Sta	atement						
	No	ovelty (N)	Yes:	Claims				



International application No. PCT/BE00/00066

No:

Yes:

Claims 1-35

Claims 1-35

Inventive step (IS)

Yes: Claims

s. Ciairii

No: Claims 1-35

No: Claims

2. Citations and explanations see separate sheet

Industrial applicability (IA)

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made: see separate sheet

EXAMINATION REPORT - SEPARATE SHEET

Re Item III

Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

Claims 1-35 were not searched in their entirety. In fact the wording of these claims abounded in unclear, vague and undefined expressions that a complete search was not deemed to be possible. What was searched were the real examples provided in the text (polymers of N-(2-hydroxyethyl)-glutamine, N-(2-hydroxypropyl)-glutamine and N-(2dihydroxypropyl)-glutamine) and pertinent methods for their preparation, their conjugates and their use for the modification of biologically active materials. Despite the statement issued with the Search Report, also claims 28-35 were searched only insofar as limited to the above subject.

Re Item V

Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

D1: WO-A-9819710 (Schacht et al.);

D2: WO-A-9736616 (University of Birmingham);

D3: CA 118: 45611 (1992).

- 1) D2 (see throughout) discloses the preparation of polymers of N-(2-hydroxyethyl)glutamine and their use for a pertinent purpose. Another example of the preparation of these polymers and of their use for pertinent purposes is provided by D3. Therefore claims 1-35 as further defined under the previous heading, are objected to under Art. 33(2) PCT because they are not new.
- 2) In the light of the (partially overlapping) documents D2 and D3, as well as of the general disclosure D1 (from the same applicant) the problem underlying the present application is set as to provide alternative polymers for the modification of biologically active materials. This scope is reached by the polymers of N-(2-hydroxypropyl)glutamine and N-(2-dihydroxypropyl)-glutamine), which were previously undisclosed

INTERNATIONAL PRELIMINARY

International application No. PCT/BE00/00066

EXAMINATION REPORT - SEPARATE SHEET

and have been demonstrated to have been prepared and also to have been used for a pertinent scope. However, these compounds, whose formula can be formulated very easily starting from polymers of the N-(2-hydroxyethyl)-glutamine, are prima facie obvious and therefore not inventive under Art. 33(3) PCT.

3) Claims 1-35 are endowed with industrial applicability under Art. 33(4) PCT.

Re Item VIII

Certain observations on the international application

Claims 1-35 are objected to under Art. 6 and Rule 6 PCT, as they contain many unclear and vague expressions; moreover, the use made by the applicant of the term "example" in the description makes completely unclear which are the real compounds prepared to embody the present application, which are the intermediate ones etc.



tional Application No.

	INTERNATIONAL SEARCH RE		00/00066		
A. CLASSI	FICATION OF SUBJECT MATTER	101/36			
IPC 7 C08G69/10 A61K47/48					
According to	o International Patent Classification (IPC) or to both national classific	ation and IPC			
	SEARCHED				
Minimum do IPC 7	ocumentation searched (classification system followed by classificati COSG A61K	on symbols)			
Documenta	tion searched other than minimum documentation to the extent that s	such documents are included in the fields	s searched		
l	lata base consulted during the international search (name of data ba BS Data, WPI Data	se and, where practical, search terms us	sed)		
C. DOCUM	ENTS CONSIDERED TO BE RELEVANT				
Category °	Citation of document, with indication, where appropriate, of the rel	evant passages	Relevant to claim No.		
Α	WO 98 19710 A (SCHACHT ET AL.) 14 May 1998 (1998-05-14) example 7		1-35		
А	WO 97 36616 A (SEYMOUR ET AL.) 9 October 1997 (1997-10-09) page 2 -page 5		1-35		
		-/			
ļ					
X Furti	her documents are listed in the continuation of box C.	X Patent family members are liste	ed in annex.		
"A" docume consid "E" earlier of filing d "L" docume which citation "O" docume other r "P" docume later th	ent defining the general state of the art which is not lered to be of particular relevance document but published on or after the international late ent which may throw doubts on priority claim(s) or is cited to establish the publication date of another n or other special reason (as specified) ent referring to an oral disclosure, use, exhibition or means ent published prior to the international filing date but an the priority date claimed	"T" later document published after the in or priority date and not in conflict will call to understand the principle or invention." "X" document of particular relevance; the cannot be considered novel or canninvolve an inventive step when the cannot be considered to involve an document of particular relevance; the cannot be considered to involve an document is combined with one or ments, such combination being obvin the art. "8" document member of the same pate	the application but theory underlying the e claimed invention to be considered to document is taken alone e claimed invention inventive step when the more other such document is underlying the consideration of the constant		
Date of the	actual completion of the international search	Date of mailing of the international s	earch report		
	9 May 2001	13/06/2001			
Name and n	nailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Masturzo, P			

3

INTERNATIONAL SEARCH REPORT

Int. dional Application No PCT/BE 00/00066

		PCT/BE 00/00066
C.(Continua	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DATABASE CHEMABS 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; DE MARRE, ANNE ET AL: "Preparation of 4-nitrophenyl carbonate esters of poly'5N-(2-hydroxyethyl) L-glutamine! and coupling with bioactive agents" retrieved from STN Database accession no. 118:45611 CA XP002168431 & MAKROMOL. CHEM. (1992), 193(12), 3023-30	1-35
Α	1992, abstract GB 781 202 A (F J WEYMOUTH) 14 August 1957 (1957-08-14)	1-35
A	the whole document K E GONSALVES & P M MUNGARA: "Synthesis and properties of degradable polyamides and related polymers" TRENDS IN POLYMER SCIENCE., vol. 4, no. 1, January 1996 (1996-01), pages 25-31, XP004049307 ELSEVIER SCIENCE PUBLISHERS B.V. AMSTERDAM., NL ISSN: 0966-4793 the whole document	1-35

International Application No. PCT/BE 00 00066

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1-27

Present claims 1-27 relate to an extremely large number of possible compounds and related methods. In fact, the claims contain so many options and variables that a lack of clarity within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible. Consequently, the search has been carried out for those parts of the application which do appear to be clear (and/or concise), namely claims 1-27 as further limited to the polymers of the poly-'N-(2-hydroxyethyl)-glutamine!, poly-'N-(2-hydroxypropyl)-glutamine! and poly-'N-(2-dihydroxypropyl)-glutamine! as in the only provided examples, as well as to the methods for their preparation, to their conjugates and their use.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.



information on patent family members

Int. Jonal Application No PCT/BE 00/00066

Patent document cited in search repo	rt	Publication date		Patent family member(s)	Publication date
WO 9819710	Α	14-05-1998	AU EP	4873997 A 0941123 A	29-05-1998 15-09-1999
WO 9736616	Α	09-10-1997	EP	0891191 A	20-01-1999
GB 781202	Α	14-08-1957	NONE		

PAILINT COOPERATION TREATY

From the INTERNATIONAL BUREAU

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

Commissioner US Department of Commerce United States Patent and Trademark

Office, PCT

2011 South Clark Place Room

CP2/5C24

Arlington, VA 22202

Date of mailing (day/month/year) 15 February 2001 (15.02.01)	ETATS-UNIS D'AMERIQUE in its capacity as elected Office		
International application No. PCT/BE00/00066	Applicant's or agent's file reference R1562-PCT		
International filing date (day/month/year) 19 June 2000 (19.06.00)	Priority date (day/month/year) 17 June 1999 (17.06.99)		
Applicant			
SCHACHT, Etienne, Honoré et al			

	The desired and Office is bounded as its placetion mode:
1.	<u> </u>
	in the demand filed with the International Preliminary Examining Authority on:
İ	10 January 2001 (10.01.01)
	in a notice effecting later election filed with the International Bureau on:
2.	The election X was
	was not
	made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).
	BEST AVAILABLE COPY
	"LABLE COPY

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland

Authorized officer

Juan Cruz

Telephone No.: (41-22) 338.83.38

Facsimile No.: (41-22) 740.14.35

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization International Bureau





(43) International Publication Date 28 December 2000 (28.12.2000)

PCT

(10) International Publication Number WO 00/78791 A2

(51) International Patent Classification⁷: C07K 1/00

(21) International Application Number: PCT/BE00/00066

(22) International Filing Date: 19 June 2000 (19.06.2000)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data: 99870125.4

17 June 1999 (17.06.1999) E

(71) Applicant (for all designated States except US): UNI-VERSITEIT GENT [BE/BE]; St. Pietersnieuwstraat 25, B-9000 Gent (BE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): SCHACHT, Etienne, Honoré [BE/BE]; Rysseveldstraat 99, B-8840 Staden (BE). TONCHEVA, Veska [BG/BE]; Pacificatielaan 5, B-9000 Gent (BE).

(74) Agents: BIRD, William et al.; Bird Goën & Co., Vilvoordsebaan 92, B-3020 Winksele (BE). (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

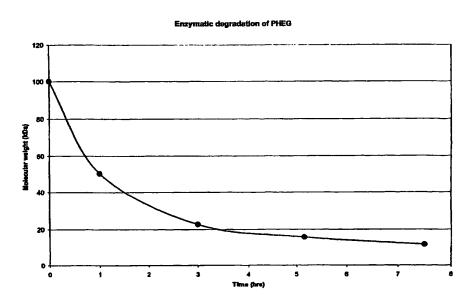
(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

 Without international search report and to be republished upon receipt of that report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: FUNCTIONAL POLY- α -AMINOACID DERIVATIVES USEFUL FOR THE MODIFICATION OF BIOLOGICALLY ACTIVE MATERIALS AND THEIR APPLICATION



(57) Abstract: A linear poly-α-amino-acid derivative has at least glutamic or aspartic or serinic repeating units and additionally having a functional group at one or both ends of the polymer backbone and/or only a single functional group as a side group on the polymer backbone, the said functional end group and/or side group being other than alcohol. The said functional derivative is useful for the modification of biologically active materials.

00/78791 A2



INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	FOR FURTHER see Notification of (Form PCT/ISA/2	of Transmittal of International Search Report (20) as well as, where applicable, item 5 below.
International application No.	International filing date (day/month/year)	(Earliest) Priority Date (day/month/year)
PCT/BE 00/00066	19/06/2000	17/06/1999
Applicant		
UNIVERSITEIT GENT		
This International Search Report has bee according to Article 18. A copy is being tr	n prepared by this International Searching Aut ansmitted to the International Bureau.	hority and is transmitted to the applicant
This International Search Report consists X It is also accompanied by	of a total of sheets. If a copy of each prior art document cited in this	s report.
Basis of the report		
With regard to the language, the language in which it was filed, un	international search was carried out on the balless otherwise indicated under this item.	sis of the international application in the
Authority (Rule 23.1(b)).	vas carried out on the basis of a translation of	
was carried out on the basis of the	e sequence listing:	nternational application, the international search
	onal application in written form. ernational application in computer readable for	m
		•••
 	o this Authority in written form. o this Authority in computer readble form.	
the statement that the su	bsequently furnished written sequence listing of as filed has been furnished.	does not go beyond the disclosure in the
		is identical to the written sequence listing has been
2. X Certain claims were for	und unsearchable (See Box I).	
3. Unity of invention is la	cking (see Box II).	
4. With regard to the title,		
the text is approved as s	ubmitted by the applicant.	
the text has been establi	shed by this Authority to read as follows:	
5. With regard to the abstract,	I with a but the combiners	
the text has been estable	submitted by the applicant. ished, according to Rule 38.2(b), by this Autho ne date of mailing of this international search re	rity as it appears in Box III. The applicant may, eport, submit comments to this Authority.
	blished with the abstract is Figure No.	
as suggested by the app		X None of the figures.
because the applicant fa		
because this figure bette	er characterizes the invention.	

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1-27

Present claims 1-27 relate to an extremely large number of possible compounds and related methods. In fact, the claims contain so many options and variables that a lack of clarity within the meaning of Article 6 PCT arises to such an extent as to render a meaningful search of the claims impossible. Consequently, the search has been carried out for those parts of the application which do appear to be clear (and/or concise), namely claims 1-27 as further limited to the polymers of the poly-'N-(2-hydroxyethyl)-glutamine!, poly-'N-(2-hydroxypropyl)-glutamine! and poly-'N-(2-dihydroxypropyl)-glutamine! as in the only provided examples, as well as to the methods for their preparation, to their conjugates and their use.

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			FC1/BE 00/00000
A. CLASSIF IPC 7	FICATION OF SUBJECT MATTER C08G69/10 A61K47/48		
According	o International Patent Classification (IPC) or to both national clas	sification and IPC	
B. FIELDS S		<u> </u>	
	ocumentation searched (classification system followed by classif	ification symbols)	
Documentati	tion searched other than minimum documentation to the extent t	that such documents are include	ded in the fields searched
Electronic da	ata base consulted during the international search (name of dat	ta base and, where practical,	search terms used)
	BS Data, WPI Data		
C. DOCUME	ENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the	he relevant passages	Relevant to claim No.
A	WO 98 19710 A (SCHACHT ET AL.) 14 May 1998 (1998-05-14) example 7)	1-35
Α	WO 97 36616 A (SEYMOUR ET AL.) 9 October 1997 (1997-10-09) page 2 -page 5)	1-35
		-/	
X Furt	orther documents are listed in the continuation of box C.	X Patent family	members are listed in annex.
Special ca 'A' docume consider 'E' earlier filing of the citation other 'P' docume other 'P' docume other	nent which may throw doubts on priority claim(s) or this cited to establish the publication date of another ion or other special reason (as specified) ment referring to an oral disclosure, use, exhibition or means ment published prior to the international filing date but	or priority date and cited to understand invention "X" document of particular cannot be considered involve an invention and the considered cannot be considered document is combined in the art.	blished after the international filing date id not in conflict with the application but he to the principle or theory underlying the cular relevance; the claimed invention ered novel or cannot be considered to investep when the document is taken alone cular relevance; the claimed invention ered to involve an inventive step when the bined with one or more other such docubination being obvious to a person skilled
later t	r than the priority date claimed le actual completion of the international search		f the international search report
	29 May 2001	13/06/2	
	d mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2	Authorized officer	
	NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Mastura	zo, P



C (Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DATABASE CHEMABS 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; DE MARRE, ANNE ET AL: "Preparation of 4-nitrophenyl carbonate esters of poly'5N-(2-hydroxyethyl) L-glutamine! and coupling with bioactive agents" retrieved from STN Database accession no. 118:45611 CA XP002168431 & MAKROMOL. CHEM. (1992), 193(12), 3023-30 '1992, abstract	1-35
Α	GB 781 202 A (F J WEYMOUTH) 14 August 1957 (1957-08-14) the whole document	1-35
A	K E GONSALVES & P M MUNGARA: "Synthesis and properties of degradable polyamides and related polymers" TRENDS IN POLYMER SCIENCE., vol. 4, no. 1, January 1996 (1996-01), pages 25-31, XP004049307 ELSEVIER SCIENCE PUBLISHERS B.V. AMSTERDAM., NL ISSN: 0966-4793 the whole document	1-35

INTERNATIONAL SEARCH REPORT

nice ation on patent family members

il ational Application No
PCT/BE 00/00066

Patent document cited in search report		Publication date	Patent family member(s)		Publication date	
WO 9819710	Α	14-05-1998	AU EP	4873997 A 0941123 A	29-05-1998 15-09-1999	
WO 9736616	Α	09-10-1997	EP	0891191 A	20-01-1999	
GB 781202	A	14-08-1957	NONE			

1/2

PATENT COOPERATION TREATMENT 2 4 BEC 2831

PCT

VAISO FOI

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

		10/00/00
Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
R1562-PCT	totamentine tille a deta (decidence	th/vear) Priority date (day/month/year)
International application No.	International filing date (day/mon	17/06/1999
PCT/BE00/00066	19/06/2000	17700/1333
International Patent Classification (IPC) or r C07K1/00	ational classification and IPC	RECEIVED
· ·		SEP 2 5 2002
Applicant UNIVERSITEIT GENT et al.		TECH CENTER 1600/290
This international preliminary examples and is transmitted to the applicant.	mination report has been prepare according to Article 36.	ed by this International Preliminary Examining Authority
2. This REPORT consists of a total of	of 5 sheets, including this cover	sheet.
heen amended and are the b	ied by ANNEXES, i.e. sheets of asis for this report and/or sheets 607 of the Administrative Instruc	the description, claims and/or drawings which have containing rectifications made before this Authority tions under the PCT).
These annexes consist of a total	of 10 sheets.	
3. This report contains indications re	elating to the following items:	
Ⅱ □ Priority		
	f opinion with regard to novelty, i	nventive step and industrial applicability
IV Lack of unity of inver		
∨ ⊠ Reasoned statement	under Article 35(2) with regard tations suporting such statement	o novelty, inventive step or industrial applicability;
VI Certain documents		
VII Certain defects in the	e international application	
	on the international application	
Date of submission of the demand	Date	of completion of this report
10/01/2001	19.12	2.2001
Name and mailing address of the international	onal Auth	prized officer
preliminary examining authority: European Patent Office - P.E NL-2280 HV Rijswijk - Pays Tel. +31 70 340 - 2040 Tx: 3	Bas Mas	sturzo, P
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INTERNATIONAL PRÉLIMINARY EXAMINATION REPORT

International application No. PCT/BE00/00066

l.	Ba	sis	of	the	re	por	t
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••		0 01 1110 10 1				ish have been furnished to		
1.	With regard to the elements of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)): Description , pages:							
	1-52		as originally filed					
	Clai	ms, No.:						
	1-36	;	as received on	06/12/2001	with letter of	26/11/2001		
	Drav	wings, sheets:						
	1/2,2	2/2	as originally filed					
2.	With regard to the language , all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.							
	These elements were available or furnished to this Authority in the following language: , which is:							
	☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).							
			publication of the internat					
		the language of a 55.2 and/or 55.3)	translation furnished fo	r the purposes of inte	rnational prelimir	nary examination (under Rule		
3.	With inte	n regard to any nu rnational prelimina	icleotide and/or amino ary examination was car	acid sequence discloring acid sequence discloring discloring the basis of the basis	osed in the intern of the sequence I	ational application, the isting:		
		contained in the i	international application	in written form.				
			n the international applic		dable form.			
	☐ furnished subsequently to this Authority in written form.							
	☐ furnished subsequently to this Authority in computer readable form.							
	☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.							
		The statement th		ded in computer reada	able form is ident	ical to the written sequence		
4	. The amendments have resulted in the cancellation of:							
		the description,	pages:					
		the claims,	Nos.:					

INTERNATIONAL PRÉLIMINARY EXAMINATION REPORT

International application No. PCT/BE00/00066

		the drawings,	sheets:		
5.		considered to go beyo	ond the dis	closure a	
		(Any replacement she report.)	eet contain	ing such	amendments must be referred to under item 1 and annexed to this
6.	Add	litional observations, if	necessary	<i>r</i> :	
V.	Rea cita	asoned statement und ations and explanatio	der Article ns suppor	: 35(2) wi rting suc	ith regard to novelty, inventive step or industrial applicability; h statement
1.	Sta	tement			
	Nov	velty (N)	Yes: No:	Claims Claims	1-36
	Inve	entive step (IS)	Yes: No:	Claims Claims	1-36
	Ind	ustrial applicability (IA)) Yes:	Claims	1-36

2. Citations and explanations see separate sheet

VIII. Certain observations on the international application

No:

Claims

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made: see separate sheet

Re Item III

Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

Claims 1-36 were now searched in their entirety.

Re Item V

Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

D1: WO-A-9819710 (Schacht et al.);

D2: WO-A-9736616 (University of Birmingham);

D3: CA 118: 45611 (1992).

- 1) D2 (see throughout) discloses the preparation of polymers of N-(2-hydroxyethyl)glutamine and their use for a pertinent purpose. Another example of the preparation of these polymers and of their use for pertinent purposes is provided by D3. Therefore claims 1-36 are objected to under Art. 33(2) PCT because they are not new. In fact all polymers of amino acids which are not specifically modified exhibit a free carboxy and a free amino group; those groups are reactive and situated at the extremities of the molecule.
- 2) In the light of the (partially overlapping) documents D2 and D3, as well as of the general disclosure D1 (from the same applicant) the problem underlying the present application is set as to provide alternative polymers for the modification of biologically active materials. This scope is reached by the some polymers of the present application which are therefore inventive under Art. 33(3) PCT. However, this cannot be extended to the ensemble of the claims, which have been objected to in the previous heading.
- 3) Claims 1-36 are endowed with industrial applicability under Art. 33(4) PCT.

Re Item VIII

Certain observations on the international application

Claims 1-36 are objected to under Art. 6 and Rule 6 PCT. In fact there are only examples of different modified poly-glutamine, whereas all polymers, also including non-amino-acid component, might fall under claim 1ff. and this makes most of the present application devoid of support.

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28. 11. 200:

1. A linear monofunctional or multifunctional poly- α -amino-acid derivative having at least glutamic or aspartic or serinic repeating units in the polymer backbone, the said glutamic or aspartic or serinic repeating units having the formula:

wherein:

- R is defined as –(CH₂)_n- CO OR₁ or –(CH₂)_n- CO NHR₂ or CH₂OH,
- n is 1 or 2,
- R_1 is selected from hydrogen, C_{1-20} alkyl, polyhalo C_{1-6} alkyl, aryl C_{1-6} alkyl and heteroaryl C_{1-6} alkyl, and
- R₂ is C₁₋₆ alkyl substituted with at least one alcohol group, characterized in additionally having a functional group at one or both ends of the polymer backbone, the said functional end group(s) being selected from amine, carboxyl, ester, carbonate, thiol, thiol precursor, thioisocyanate, thiocarbonate, urea, thiourea, aldehyde, acetal, N-carboxyanhydride, oxycarbonyl, maleimide or any vinyl group suitable for radical, anionic or cationic polymerization.
- 2. A linear multifunctional poly- α -amino-acid derivative according to claim 1, having a functional group at both ends of the polymer backbone, characterized in additionally having a single functional group as a side group.
- 3. A linear poly-α-amino-acid derivative according to claim 2, wherein the said functional side group is selected from the group consisting of amine, thiol precursor, thioisocyanate, thiocarbonate, urea, thiourea, acetal, N-carboxyanhydride, oxycarbonyl, maleimide and any vinyl group suitable for radical, anionic or cationic polymerization.
- 4. A linear multifunctional poly- α -amino-acid derivative according to claim 2, wherein the said functional side group is selected from carboxyl,

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ester, carbonate, thiol and aldehyde.

- 5. A linear poly- α -amino-acid derivative according to any of claims 1 to 4, additionally comprising repeating units of one or more comonomer(s) copolymerizable with the α -amino-acid sequence containing glutamic or aspartic or serinic repeating units.
- 6. A linear poly- α -amino-acid derivative according to claim 5, wherein the said co-monomer is any naturally occurring α -amino-acid other than glutamic acid, aspartic acid and serine.
- 7. A linear poly- α -amino-acid derivative according to claim 5, wherein the said co-monomer is a polymer block or sequence derived from ethylene oxide or propylene oxide or mixtures thereof or from a polyhydroxyalkanoate.
- 8. A linear poly- α -amino-acid derivative according to any of claims 1 to 7, being multifunctional and having any of the following formulae:

$$V-[CO-CHR-NH]_{x^{-}}[A]_{y^{-}}W$$
 (IIa)

$$V-[A]_{y^{-}}[CO-CHR-NH]_{x^{-}}W$$
 (IIb)

$$V-[CO - CHR - NH]_{x^{-}}[A]_{y^{-}}[CO - CHR - NH]_{x^{-}}W$$
 (IIc)

$$V-[CO-CHR-NH]_{x^{-}}T-[CO-CHR-NH]_{x^{\prime -}}V \hspace{1cm} (IId)$$

W

V-[CO - CHR - NH]_x- T -[CO - CHR - NH]_{x'}- V' (IIe)
$$W$$

wherein:

- R is as defined in claim 1,
- x or, where applicable, x + x' range from 2 to 2,000,
- each of V and W independently represent a functional group,
- A is at least a co-monomer copolymerizable with the α -amino-acid sequence containing glutamic or aspartic or serinic repeating units,
- y ranges from 0 to 500,
- T is a spacing unit selected from lysine and ornithine, and

- V' is a non-reactive end group.
- 9. A linear poly- α -amino-acid derivative according to any of claims 1 to 7, being monofunctional and having any of the following formulae:

$$V-[CO - CHR - NH]_{x}-[A]_{y}-W'$$

(Va)

(Vb)

$$V-[CO - CHR - NH]_{x^{-}}[A]_{y^{-}}[CO - CHR - NH]_{x'}-W'$$
 (Vc)

$$V'-[CO-CHR-NH]_{x^{-}}[A]_{y^{-}}[CO-CHR-NH]_{x^{-}}W$$
 (Vd)

$$V'-[CO - CHR - NH]_{x}-T-[CO - CHR - NH]_{x}-V'$$
 (VI)

W

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wherein:

- R is as defined in claim 1,
- x or, where applicable, x + x' range from 2 to 2,000, and
- each of V and W independently represent a functional group,
- A is at least a co-monomer copolymerizable with the α -amino-acid sequence containing glutamic or aspartic or serinic repeating units,
- y ranges from 0 to 500,
- T is a spacing unit selected from lysine and ornithine, and
- V' and W' are non-reactive end groups.

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10. A linear poly-α-amino-acid derivative according to claim 8 or claim 9, wherein A is represented by the formula - CO - CHR' - NH - (III), wherein R' is the side-chain group of an α-amino acid other than glutamic acid or aspartic acid or serine, or by the formula

 $CH_2 - CHR'' - X' - (IV)$, wherein:

- R" is selected from hydrogen and methyl, and
- X' is selected from a single bond and oxygen,
 or A is a repeating unit derived from a hydroxyalkanoate.
- 11. A linear poly-α-amino-acid derivative according to any of claims 8 to 10, wherein each of V' and/or W' is selected from C₁₋₂₀ alkyl, aryl, amide,oxyC₁₋₂₀alkyl, arylC₁₋₂₀ alkyl, heteroaryl and heteroarylC₁₋₂₀ alkyl.

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12. A linear poly- α -amino-acid derivative according to claim 1, having at least one protective end group and being represented by the following formulae:

$$X_{\Gamma}NH$$
 $= \begin{bmatrix} O & O & O \\ C & -CH - NH \end{bmatrix}_{X} = \begin{bmatrix} O & C & CH - NH \end{bmatrix}_{X} = \begin{bmatrix} O & CH - NH$

$$Y_2 - R_3 - NH$$

$$\begin{array}{c} O \\ \parallel \\ C - CH - NH \\ \parallel \\ X \end{array}$$

$$\begin{array}{c} O \\ \parallel \\ X \end{array}$$

$$\begin{array}{c} C \\ \downarrow \\ X_2 \end{array}$$

$$\begin{array}{c} (VIII), \end{array}$$

wherein:

- R is $-(CH_2)_n$ CO NHR_2 ,
- R2and n are as defined in claim 1,
- x ranges from 2 to 2,000,

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$$X_1$$
 is - R_4 - Z_1 - A_1 ,

- each of R_3 and R_4 is independently selected from $(CH_2)_m$, arylene, C_{1-6} alkylarylene and aryl C_{1-6} alkylene,
- m is from 2 to 20,

$$-Y_1$$
 is $-Z_2 - A_2$,

-
$$X_2$$
 is - R_4 - Z_3 - A_3 or - $O - R_4$ - Z_3 - A_3 ,

$$-Y_2$$
 is $-Z_4 - A_4$,

- each of Z_1 , Z_2 , Z_3 and Z_4 is independently selected from NH, O, S, C(O)O, C(S)O, CO, CS, -OCH-O- and C = N R₅,
- each of A_1 , A_2 , A_3 and A_4 is a protective group suitable for Z_1 , Z_2 , Z_3 and Z_4 respectively, and
- R_5 is selected from hydrogen, $C_{1\text{-}6}$ alkyl, aryl and $C_{1\text{-}6}$ alkylaryl, heteroaryl and $C_{1\text{-}6}$ alkylheteroaryl.
- 13. A linear poly- α -amino-acid derivative according to claim 1, being represented by the formula:

$$X_1$$
-NH-[CO - CHR - NH]_x- CO - CHR - NH₂ (IX)

wherein:

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$$X_1$$
 is $-R_4 - Z_1 - A_1$,

- R₄ is selected from (CH₂)_m, arylene, C₁₋₆ alkylarylene and arylC₁₋₆

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alkylene,

- x ranges from 2 to 2,000,
- R is defined as -(CH₂)_n- CO OR₁,
- R₁ and n are as defined in claim 1,
- Z_1 is selected from NH, O, S, C(O)O, C(S)O, CO, CS, -OCH-O- and $C = N R_5$,
 - A₁ is a protective group suitable for Z₁, and
 - R_5 is selected from hydrogen, $C_{1\text{-}6}$ alkyl, aryl and $C_{1\text{-}6}$ alkylaryl, heteroaryl and $C_{1\text{-}6}$ alkylheteroaryl.
 - 14. A linear poly- α -amino-acid derivative according to claim 1, being represented by any of the respective formulae:

$$D_{4}$$
 R_{3} $-NH + C - CH - NH + C - X_{2}$

(X), and

$$X_{\Gamma}NH$$

$$\begin{array}{c}
O \\
H \\
C \\
C \\
R
\end{array}$$

$$\begin{array}{c}
O \\
H \\
C \\
R
\end{array}$$

$$\begin{array}{c}
O \\
H \\
C \\
R_3 \\
\end{array}$$

$$\begin{array}{c}
Z_2 \\
R_3 \\
\end{array}$$

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(XI), wherein:

- R is $-(CH_2)_n$ CO NHR_2 ,
- R₂ and n are as defined in claim 1,
- x ranges from 2 to 2,000;
- X_1 is R_4 Z_1 D_1 ,

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- each of R_3 and R_4 is independently selected from $(CH_2)_m$, arylene, C_{1-6} alkylarylene and aryl C_{1-6} alkylene,
- m is from 2 to 20,
- each of $R_3 Y_1$ and $R_3 Y_2$ may be a group including a vinyl terminal moiety,

- $-X_2$ is $-R_4-Z_3-D_3$,
- each of Z_1 , Z_2 , Z_3 and Z_4 is independently selected from NH, O, S, C(O)O, C(S)O, CO, CS, -OCH-O- and C = N R₅,

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- each of D_1 , D_2 , D_3 and D_4 is independently selected from hydrogen, aryl, heteroaryl, succinimidyl, vinyl, C_{1-6} alkylcarbonyl,
- each of Z_1 D_1 , Z_2 D_2 , Z_3 D_3 and Z_4 D_4 may be independently selected from maleimidyl, disulfide, α -haloacetoxy and C_{1-6} alkyloxy-methylsulfide, and
- R_5 is selected from hydrogen, C_{1-6} alkyl, aryl and C_{1-6} alkylaryl, heteroaryl and C_{1-6} alkylheteroaryl.
- 15. A linear poly- α -amino-acid derivative according to claim 14, wherein D_1 is different from D_2 and D_3 is different from D_4 .
 - 16. A process for making a linear poly-α-amino-acid derivative according to any of claims 1 to 15, including a step comprising polymerizing a monomer or mixture of monomers comprising at least the N-carboxy anhydride of an amino-acid selected from glutamic acid, aspartic acid, serine and oxygen-protected serine in the presence of an effective amount of a multifunctional initiator containing at least one primary amino group and further containing at least another functional group selected from maleimide, thioisocyanate, thiocarbonate, urea, thiourea, aldehyde, acetal, oxycarbonyl, vinyl, ester, carbonate, thiol precursor, protected amine and protected carboxylic acid and/or in the presence of an effective amount of a bi-functional terminating reagent.
 - 17. A process according to claim 16, wherein the multifunctional initiator is selected from amino-acid esters, α-amino-ω-diC₁₋₆alkylacetals, α,α' -diamino C₁₋₆alkyldisulfides and α-amino-ω-maleimido alkanoic acid amides.
 - 18. A process according to claim 16 or claim 17, wherein the amount of the multifunctional initiator ranges between 0.2 and 30 mole % with respect to the N-carboxy-anhydride monomer.
 - 19. A process according to any of claims 16 to 18, wherein the amount of

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the bi-functional terminating reagent ranges between 2 and 5 equivalents with respect to the molar amount of the multifunctional initiator.

- 20. A process according to any of claims 16 to 19, further including aminolysis of the pending R₁ group of the glutamic, aspartic or serinic repeating unit derived from glutamic acid, aspartic acid or serine by means of an effective amount of an amino-alcohol, in the presence of an effective amount of a reaction promoter.
- 21. A process according to claim 20, wherein the effective amount of the amino-alcohol used during the said aminolysis step ranges from 1 to 50, equivalents with respect to the monomeric units in the polymer.
- 22. A process according to claim 20 or claim 21, wherein the effective amount of the reaction promoter ranges from 0.5 to 5 equivalents with respect to the monomeric units in the polymer.
 - 23. A process for making a linear poly-α-amino-acid derivative according to any of claims 1 to 15, including:
 - a first step of N-acylating part of an α -amino-acid selected from glutamic acid, aspartic acid and serine, then separately treating the N -acylated α -amino-acid and the remaining part of the said α -amino-acid in order to form a mixture of the corresponding N-carboxy anhydrides, and
 - a second step of copolymerizing the said mixture of N-carboxy anhydrides in the presence of an initiator.
 - 24. A process according to claim 23, wherein the N-carboxy anhydride of the α -amino-acid is used in excess of the N-carboxy anhydride of the N-acylated α -amino-acid.
 - 25. A process according to claim 23 or claim 24, wherein the N-carboxy

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anhydride-terminated polymer obtained in the second step is reacted with a reagent having the formula $H_2N-R_3-Y_2$, wherein:

- R_3 is selected from $(CH_2)_m$, arylene, C_{1-6} alkylarylene and aryl C_{1-6} alkylene,
- Y_2 is $-Z_4 A_4$,
- Z₄ is selected from NH, O, S, C(O)O, C(S)O, CO, CS, -OCH-O- and
 C = N R₅,
- A₄ is a protective group suitable for Z₄, and
- R_5 is selected from hydrogen, C_{1-6} alkyl, aryl and C_{1-6} alkylaryl, heteroaryl and C_{1-6} alkylheteroaryl.
- 26. A biodegradable article containing a copolymer comprising at least a moiety derived from a poly- α -amino-acid derivative according to any of claims 1 to 15, provided that the functional group at one or both ends thereof is an unsaturated group.
- 27. Use of a poly- α -amino-acid derivative according to any of claims 1 to 15 for the modification of a biologically-active ingredient.
- 28. An enzymatically degradable poly-α-amino-acid derivative according to any of claims 1 to 15, containing a L-amino-acid sequence.
 - 29. The product of coupling a poly- α -amino-acid derivative according to any of claims 1 to 15 with a biomolecule.
 - 30. The product of claim 29, wherein the said biomolecule is a therapeutic agent, prophylactic agent, diagnostic agent, protein, peptide, hormone, antibody or fragment thereof, oligonucleotide, plasmid, DNA, interleukin, interferon, enzyme or fragment thereof.
 - 31. The product of claim 29 or claim 30, being an antibody modified by means of the said functional poly-α-aminoacid derivatives and having a second functionality for hooking and/or being able to attach

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another targeting group such as an antibody, a peptide, an oligopeptide or a saccharide.

- 32. Use of a non degradable poly- α -amino-acid derivative according to any of claims 1 to 15, containing a D-amino-acid sequence, for the surface modification of a biomaterial.
- 33. A synthetic polymer for a polymer-based carrier vehicle or vector for delivery of DNA or other nucleic acid material to target cells in a biological system, comprising a linear poly-α-amino-acid derivative according to any of claims 1 to 15.
- 34. A synthetic polymer for a polymer-based carrier vehicle or vector according to claim 33, further comprising a synthetic vector component such as polyethyleneimine, poly-L-lysine, a star-shaped dendrimer or chitosan.
- 35. A method of treatment of a patient in need of such treatment, comprising administration to said patient of a biologically-active ingredient modified by or a nucleic acid material carried by a polymer system comprising a linear poly-α-amino-acid derivative according to any of claims 1 to 15.
- 36. A linear monofunctional or multifunctional poly-α-amino-acid derivative having at least glutamic or aspartic or serinic repeating units in the polymer backbone, the said glutamic or aspartic or serinic repeating units having the formula:

$$-CO-CHR-NH-$$
 (1)

wherein:

- R is defined as –(CH₂)_n- CO OR₁ or –(CH₂)_n- CO NHR₂ or CH₂OH,
- n is 1 or 2,
- R₁ is selected from hydrogen, C₁₋₂₀ alkyl, polyhaloC₁₋₆alkyl, arylC₁₋₆



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alkyl and heteroarylC₁₋₆ alkyl, and

- R_2 is C_{1-6} alkyl substituted with at least one alcohol group, characterized in additionally having a functional group at one or both ends of the polymer backbone, the said functional end group(s) being other than alcohol.

PATENT COOPERATION TREATY

RECT 2.4 DEC 2001

- From the INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

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1.		demand (Rule 61.1(a)).			
2.	×	copy of the international preliminary examination report and its annexes (Rule 71.1).			
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